FILE 'USPAT' ENTERED AT 08:52:25 ON 29 APR 1999 WELCOME TO THE U.S. PATENT TEXT FILE => s bmp(p)(bone or protein) 677 BMP 34554 BONE 70917 PROTEIN 361 BMP(P)(BONE OR PROTEIN) => s bone(w)(morphogenetic or morphogenic)(w)protein 34554 BONE **481 MORPHOGENETIC 413 MORPHOGENIC** 70917 PROTEIN 407 BONE(W)(MORPHOGENETIC OR MORPHOGENIC)(W)PROTEIN => s plasmid or plasmids or vector or vectors or recombinant or recombinants 15090 PLASMID 11330 PLASMIDS **68287 VECTOR** 37880 VECTORS 20292 RECOMBINANT 2730 RECOMBINANTS 82728 PLASMID OR PLASMIDS OR VECTOR OR VECTORS OR RECOMBINANT OR **OMBINANTS** => s 11 or 12 503 L1 OR L2 => s l4(10a)l3

WARNING - PROXIMITY OPERATOR PRECEDENCE LEVEL CONFLICTS OR IS NOT CONSIS TENT WITH FIELD CODE - 'AND' OPERATOR ASSUMED 'L4(10A)L3' 289 L4(10A)L3

=> s therapy or therapies or therapeutic or treat or treats or treated or treatment or treatments or implant or implants or implantation or implanted

5859 THERAPIES 61471 THERAPEUTIC 73573 TREAT 7941 TREATS **383604 TREATED** 400754 TREATMENT 70723 TREATMENTS 21425 IMPLANT 13006 IMPLANTS 35766 IMPLANTATION 37521 IMPLANTED 600838 THERAPY OR THERAPIES OR THERAPEUTIC OR TREAT OR TREATS OR T

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43205 THERAPY

=> s 15(10a)16

WARNING - PROXIMITY OPERATOR PRECEDENCE LEVEL CONFLICTS OR IS NOT CONSIS TENT WITH FIELD CODE - 'AND' OPERATOR ASSUMED 'L5(10A)L6' 285 L5(10A)L6

=> d his

(FILE 'USPAT' ENTERED AT 08:52:25 ON 29 APR 1999) 361 S BMP(P)(BONE OR PROTEIN) 407 S BONE(W)(MORPHOGENETIC OR MORPHOGENIC)(W)PROTEIN 82728 S PLASMID OR PLASMIDS OR VECTOR OR VECTORS OR RECOMBINANT OR 503 S L1 OR L2 L4 289 S L4(10A)L3 1.5 600838 S THERAPY OR THERAPIES OR THERAPEUTIC OR 1.6 TREAT OR TREATS O 285 S L5(10A)L6

222 L4(P)L6

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WARNING - PROXIMITY OPERATOR PRECEDENCE LEVEL CONFLICTS OR IS NOT CONSIS TENT WITH FIELD CODE - 'AND' OPERATOR ASSUMED 'L4(10A)L3' 289 L4(10A)L3

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WARNING - PROXIMITY OPERATOR PRECEDENCE LEVEL CONFLICTS OR IS NOT CONSIS TENT WITH FIELD CODE - 'AND' OPERATOR ASSUMED 'L9(P)L8' LIO 119 L9(P)L8

=> d110,cit,rel,ab,1-119

1. 5,891,147, Apr. 6, 1999, Minimally invasive spinal surgical methods & instruments, Peter A. Moskovitz, et al., 606/79 [IMAGE AVAILABLE]

US PAT NO: 5,891,147 [IMAGE AVAILABLE] REL-US-DATA: Division of Ser. No. 670,351, Jun. 25, 1996, Pat. No. 5,741,261.

ABSTRACT:

Minimally invasive spinal surgical techniques and tools are provided. The methods include separating the iliocostalis lumborum muscle from the anterior leaf of the thoracolumbar fascia to create a channel from the patient's skin to the intertransverse interval. In one embodiment, the method also includes delivering graft material through the channel to the intertransverse interval. A device according to one aspect of the present invention includes a retraction portion having a flattened plate configured to atraumatically retract tissue to create a working space within an endosurgical site and a curved shaft attached to the retraction portion. The shaft includes a bend having a radius of preferably 160 degrees. A gripping portion is attached to the shaft and is configured for manually gripping and manipulating the device.

2. 5,885,829, Mar. 23, 1999, Engineering oral tissues; David J. Mooney, et al., 435/325; 424/49, 422, 435; 435/69.1, 374, 378 [IMAGE AVAILABLE]

US PAT NO: 5,885,829 [IMAGE AVAILABLE] L10: 2 of 119

Disclosed are methods for regenerating dental and oral tissues from viable cells using ex vivo culture on a structural matrix. The regenerated oral tissues and tissue-matrix preparations thus provided have both clinical applications in dentistry and oral medicine and are also useful in in vitro toxicity and biocompatibility testing.

3. 5,885,292, Mar. 23, 1999, Minimally invasive spinal surgical methods and instruments; Peter A. Moskovitz, et al., 606/79, 61, 86 [IMAGE AVAILABLE]

US PAT NO: 5,885,292 [IMAGE AVAILABLE] REL-US-DATA: Division of Ser. No. 670,351, Jun. 25, 1996, Pat. No.

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     842062 BONE
    3253189 PROTEIN
   S1 2529 BMP(10N)(BONE OR PROTEIN)
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     842062 BONE
      17821 MORPHOGENETIC
      2564 MORPHOGENIC
    3253189 PROTEIN
   S2 5309 BONE(W)(MORPHOGENETIC OR
MORPHOGENIC)(W)PROTEIN
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Set Items Description

85116 IMPLANTS

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IMPLANT OR

5355 IMPLANTING

S8 7285544 THERAPY OR THERAPIES OR THERAPEUTIC OR

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DIALOG(R)File 5:Biosis Previews(R)
(c) 1999 BIOSIS. All rts. reserv.
11895001 BIOSIS NO.: 199900141110
Effective doses of recombinant human bone morphogenetic protein-2 in
 experimental spinal fusion.
AUTHOR: Sandhu Harvinder S(a); Kanim Linda E A; Kabo J Michael; Toth
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Jeffrey M; Zeegen Erik N; Liu David; Delamarter Rick B; Dawson Edgar G AUTHOR ADDRESS: (a)UCLA Sch. Med., 200 UCLA Med. Plaza, Suite 140, Los Angeles, CA 90095-6902, USA

JOURNAL: Spine 21 (18):p2115-2122 Sept. 15, 1996 ISSN: 0362-2436 DOCUMENT TYPE: Article RECORD TYPE: Abstract LANGUAGE: English

ABSTRACT: Study Design. Nineteen dogs underwent L4-L5 intertransverse process fusions with either 58 mug, 115 mug, 230 mug, 460 mug, or 920 mug of recombinant human bone morphogenetic protein-2 carried by a polylactic acid polymer. A previous study (12 dogs) compared 2300 mug of recombinant human bone morphogenetic protein-2, autogenous iliac bone, and carrier alone in this model. All fusions subsequently were compared. Objectives. To characterize the dose-response relationship of recombinant human bone morphogenetic protein-2 in a spinal fusion model. Summary of Background Data. Recombinant osteoinductive morphogens, such as recombinant human bone morphogenetic protein-2, are effective in vertebrate diaphyseal defect and spinal fusion models. It is hypothesized that the quality of spinal fusion produced with %%%recombinant%%% human %%%bone%%%%

%%%morphogenetic%%%%%%protein%%%-2, above a threshold dose, does not

change with increasing amounts of inductive protein. Methods. After decortication of the posterior elements, the designated %%%implants%%% were placed along the intertransverse process space bilaterally. The fusion sites were evaluated after 3 months by computed tomography imaging, high-resolution radiography, manual testing, mechanical testing, and histologic analysis. Results. As in the study using 2300 mug of recombinant human bone morphogenetic protein-2, implantation of 58-920 mug of recombinant human bone morphogenetic protein-2 successfully resulted in intertransverse process fusion in the dog by 3 months. This had not occurred in animals containing autograft or carrier alone. The cross-sectional area of the fusion mass and mechanical stiffness of the L4-L5 intersegment were not dose-dependent. Histologic findings varied but were not related to rhBMP-2 dose. Inflammatory reaction to the composite implant was proportional inversely to the volume of the fusion mass. Conclusions. No mechanical, radiographic, or histologic differences in the quality of intertransverse process fusion resulted from a 40-fold variation in dose of recombinant human bone morphogenetic protein-2.

11/3,AB/2 (Item 2 from file: 5)